## II. CLAIM AMENDMENTS

1. (Currently Amended) A method for of treating a migraine  $\frac{1}{2}$  subject in need thereof comprising administering to of to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted  $\gamma$ -lactone compound of the general formula I,

in which

R<sup>1</sup> denotes an optionally at least mono-substituted 2- pyridyl, 2-pyrimidyl, 2-pyrazolyl 3-pyrazolyl, 2-quinolinyl or 2- pyrazinyl residue, which may also be fused with a saturated or at least partially unsaturated hydrocarbon ring system,

- $R^2$  denotes an optionally at least mono-substituted, saturated, branched or unbranched aliphatic  $C_{1-10}$  residue or an optionally at least mono-substituted, at least partially unsaturated, branched or unbranched aliphatic  $C_{2-10}$  residue,
- R<sup>3</sup> denotes an optionally at least mono-substituted aryl residue,
- R4 denotes H,

or

- $R^3$  and  $R^4$  together denote an optionally at least monosubstituted, saturated or at least mono-unsaturated aliphatic  $C_{3-7}$  residue, with the proviso that the residue  $R^2$  in this case denotes an optionally at least mono-substituted aryl residue, an optionally at least mono-substituted, saturated, branched or unbranched aliphatic  $C_{1-10}$  residue or an optionally at least mono-substituted, at least partially unsaturated, branched or unbranched aliphatic  $C_{2-10}$  residue,
- said compound being in the form of the a racemates, a diastereomers or enantiomers thereof as a free base or of a corresponding physiologically acceptable salt therof.for the production of a pharmaceutical preparation for the treatment of migraine.
- 2. (Currently Amended) The A method Use according to claim 1, characterised in that R<sup>1</sup> denotes an optionally at least monosubstituted 2— pyridyl-residue, which may also be fused with a saturated or at least partially unsaturated hydrocarbon ring

system, preferably denotes a 2- pyridyl residue which is substituted at least in position 4.

- 3. (Currently Amended) The A method Use according to claim 1, characterised in that  $R^2$  denotes an optionally at least monosubstituted, saturated, branched or unbranched aliphatic  $C_{1-6}$  residue.
- 4. (Currently Amended)  $\frac{\text{The A method Use}}{\text{Use}}$  according to claim 1, characterised in that  $R^3$  denotes an optionally at least monosubstituted aryl residue and  $R^4$  denotes H.
- 5. (Currently Amended) The A method Use according to claim 1, characterised in that the compound used of the general formula I is according to claim 1 comprises at least one compound selected from the group consisting of
  - 5-(2,4-Dimethyl-phenyl)-3-(8-hydroxy-quinolin-2-ylamino)-5-methyl-dihydro-furan-2-one,
  - 5-(3,4-Dimethyl-phenyl)-3-(4,6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
  - 5-(2,4-Dimethyl-phenyl)-3-(4,6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
  - 5-(4-Cyclohexyl-phenyl)-3-(4,6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,

- 5-(3,5-Dimethyl-phenyl)-3-(4,6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
- 5-(3,4-Dimethyl-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(2,4-Dimethyl-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(4-Cyclohexyl-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-Methyl-3-(quinolin-2-ylamino)-5-m-tolyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-methyl-5-p-tolyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-methyl-5-m-tolyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-(4-ethoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 4-[4-(3-Bromo-5-methyl-pyridin-2-ylamino)-2-methyl-5-oxo-tetrahydro-furan-2-yl]-benzonitrile,
- 3-(3-Bromo-5-methyl-pyridin-2-ylamino)-5-(4-tert-butyl-phenyl)-5-methyl-dihydro-furan-2-one,
- 5-(4-tert-Butyl-phenyl)-5-methyl-3-(6-propyl-pyridin-2-ylamino)-dihydro-furan-2-one,

- 5-(4-tert-Butyl-phenyl)-5-methyl-3-(4-methyl-3-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(5-Bromo-6-methyl-pyridin-2-ylamino)-5-methyl-5-(4-phenoxy-phenyl)-dihydro-furan-2-one,
- 5-(4-tert-Butyl-phenyl)-5-methyl-3-(3-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(3-Benzyloxy-pyridin-2-ylamino)-5-methyl-5-(4-phenoxy-phenyl)-dihydro-furan-2-one,
- 3—(3—Benzyloxy-pyridin-2-ylamino)-5-(4-tert-butyl-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-methyl-5-(4-phenoxy-phenyl)-dihydro-furan-2-one,
- 5-(4-tert-Butyl-phenyl)-3-(4, 6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydrofuran-2-one,
- 5-Methyl-3- (4-methyl-pyridin-2-ylamino)-5-(4-phenoxy-phenyl)-dihydro-furan-2-one,
- 5-(4-tert-Butyl-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 4-[4-(5-Bromo-3-nitro-pyridin-2-ylamino)-2-methyl-5-oxo-tetrahydro-furan-2-yl]-benzonitrile,

- 4-[4-(5-Bromo-pyrimidin-2-ylamino)-2-methyl-5-oxotetrahydro-furan-2-yl]-benzonitrile,
- 5-Benzo[b] thiophen-2-yl-5-methyl-3-(6-propyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-(4-isopropyl-phenyl)-5-methyl-dihydro-furan-2-one,
- 5-Benzofuran-2-yl-3-(4,6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
- 5-Benzo(b) thiophen-2-yl-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-Benzofuran-2-yl-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(5-Benzo[1,3]dioxol-5-yl-5-methyl-2-oxo- tetrahydro-furan-3-ylamino)-lH-pyrazole-4- carbonitrile,
- 3-(5-Benzo[1,3]dioxol-5-yl-5-methyl-2-oxo-tetrahydro-furan-3-ylamino)-1H-pyrazole-4-carboxylic acid ethyl ester,
- 5-Benzo[1, 3]dioxol-5-yl-5-methyl-3-(3-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-methyl-5-(5,6,7,8-tetrahydro-naphthalen-2-yl)-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-methyl-5-naphthalen-2-yl-dihydro-furan-2-one,

5-Benzo[1,3]dioxol-5-yl-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,

5-Methyl-3-(4-methyl-pyridin-2-ylamino)-5-(5,6,7,8-tetrahydro-naphthalen-2-yl)-dihydro-furan-2-one,

5-Benzo[1,3]dioxol-5-yl-5-methyl-3-(5-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,

5-Benzo[1,3]dioxol-5-yl-5-methyl-3-(6-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,

3-(5-Bromo-3-nitro-pyridin-2-ylamino)-5-methyl-5-(5,6,7,8-tetrahydro-naphthalen-2-yl)-dihydro-furan-2-one,

3-(5-Bromo-3-nitro-pyridin-2-ylamino)-5-isopropyl-5-phenyl-dihydro-furan-2-one,

5-Isopropyl-3-(5-nitro-pyridin-2-ylamino)-5-phenyl-dihydro-furan-2-one,

5-Methyl-5-naphthalen-2-yl-3-(5-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,

5-Isopropyl-5-phenyl-3-(pyrimidin-2-ylamino)-dihydrofuran-2-one,

3-[5-(4-Iodo-phenyl)-5-methyl-2-oxo-tetrahydro-furan-3-ylamino]-1H-pyrazole-4-carboxylic acid ethyl ester,

- 5-(4-Bromo-phenyl)-3-(5-bromo-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
- 3-(3-Bromo-5-methyl-pyridin-2-ylamino)-5-methyl-5-naphthalen-1-yl-dihydro-furan-2-one,
- 5-Methyl-5-naphthalen-1-yl-3-(6-propyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(3-Chloro-phenyl)-5-methyl-3-(6-propyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(3-Chloro-phenyl)-5-methyl-3-(4-methyl-3-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(4-Bromo-phenyl)-5-methyl-3-(4-methyl-3-nitro-pyridin-2-ylamino)-dihydro-furan-2-one,
- 3-(5-Bromo-6-methyl-pyridin-2-ylamino)-5-methyl-5-naphthalen-1-yl-dihydro-furan-2-one,
- 3-(5-Bromo-6-methyl-pyridin-2-ylamino)-5-(4-iodo-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(3-Benzyloxy-pyridin-2-ylamino)-5-(4-iodo-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(3-Benzyloxy-pyridin-2-ylamino)-5-(4-bromo-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-(4-iodo-phenyl)-5-methyl-dihydro-furan-2-one,

- 5-(3-Chloro-phenyl)-3-(4, 6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
- 5-(4-Bromo-phenyl)-5-methyl-3-(3-methyl- pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(4-Bromo-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 2-[5-(3,5-Dimethoxy-phenyl)-5-methyl-2-oxo-tetrahydro-furan-3-ylamino]-4-propyl-pyrimidine-5-carboxylic acid ethyl ester,
- 3-(4-Bromo-1H-pyrazol-3-ylamino)-5-(3,5-dimethoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(4-Bromo-1H-pyrazol-3-ylamino)-5-(2-methoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-[5-(2,5-Dimethoxy-phenyl)-5-methyl-2-oxo-tetrahydro-furan-3-ylamino]-1H-pyrazole-4-carbonitrile,
- 3-[5-(2,5-Dimethoxy-phenyl)-5-methyl-2-oxo-tetrahydro-furan-3-ylamino]-5-methylsulfanyl-1H-pyrazole-4-carbonitrile,
- 5-(2,5-Dimethoxy-phenyl)-5-methyl-3-(pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(2-Methoxy-phenyl)-5-methyl-3-(pyridin-2-ylamino)-dihydro-furan-2-one,

- 3-(3-Chloro-5-trifluoromethyl-pyridin-2-ylamino)-5-(3,5-dimethoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(3-Chloro-5-trifluoromethyl-pyridin-2-ylamino)-5-(2,5-dimethoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(3,5-Dichloro-pyridin-2-ylamino)-5-(2-methoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(3-Chloro-5-trifluoromethyl-pyridin-2-ylamino)-5-(2,4-dimethoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-(3-methoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 3-(4,6-Dimethyl-pyridin-2-ylamino)-5-(4-methoxy-phenyl)-5-methyl-dihydro-furan-2-one,
- 5-(3, 4-Dimethoxy-phenyl)-3-(4, 6-dimethyl-pyridin-2-ylamino)-5-methyl-dihydro-furan-2-one,
- 5-(4-Methoxy-phenyl)-5-methyl-3-(4-methyl-pyridin-2-ylamino)-dihydro-furan-2-one,
- 5-(2,5-Dimethoxy-phenyl)-5-methyl-3-(pyrazin-2-ylamino)-dihydro-furan-2-one and
- 5-Methylsulfanyl-3-(2-oxo-5-phenyl-5-propyl-tetrahydro-furan-3-ylamino)-1H-pyrazole-4-carbonitrile
- and the corresponding physiologically acceptable salts thereof, preferably the hydrochlorides thereof.

- 6. (Currently Amended)—The A method according to claim—I Use of using—of treating septic shock comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim—I according to claim—I for the production of a pharmaceutical preparation for the treatment of septic shock.
- 7. (Currently Amended) The A method according to claim 1 Use of using of treating a neurodegenerative disease comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted  $\gamma$ -lactone compound of the general formula I according to claim 1.according to claim 1 for the production of a pharmaceutical preparation for the treatment of neurodegenerative diseases.
- 8. (Currently Amended) The A method Use according to claim 7, wherein the neurodegenerative disease is multiple sclerosis. for the production of a pharmaceutical preparation for the treatment of multiple sclerosis.
- 9. (Currently Amended) The A method Use according to claim 7, wherein the neurodegenerative disease is Parkison's disease. for the production of a pharmaceutical preparation for the treatment of Parkinson's disease.
- 10. (Currently Amended) <u>The-A method</u> Use according to claim 7, for the production of a pharmaceutical preparation for the

treatment of wherein the neurodegenerative disease is Alzheimer's disease.

- 11. (Currently Amended) The A method Use according to claim 7, for the production of a pharmaceutical preparation for the treatment of wherein the neurodegenerative disease is Huntington's chorea.
- 12. (Currently Amended) The A method according to claim 1 Use of using of treating inflammation comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim 1.according to claim 1 for the production of a pharmaceutical preparation for the treatment of inflammation.
- 13. (Currently Amended) The-A method according to claim 1 Use of using of treating inflammatory pain comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim 1 according to claim 1 for the production of a pharmaceutical preparation for the treatment of inflammatory pain.
- 14. (Currently Amended) The A method according to claim 1 Use of using of treating cerebral ischaemia comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted  $\gamma$ -lactone

compound of the general formula I according to claim 1.according to claim 1 for the production of a pharmaceutical preparation for the treatment of cerebral ischaemia.

- 15. (Currently Amended) The A method according to claim 1 Use of using of treating diabetes comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim 1 according to claim 1 for the production of a pharmaceutical preparation for the treatment of diabetes.
- 16. (Currently Amended) The A method according to claim 1 Use of using of treating meningitis comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim 1.according to claim 1 for the production of a pharmaceutical preparation for the treatment of meningitis.
- 17. (Currently Amended) The A method according to claim 1 Use of using—of treating arteriosclerosis comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted  $\gamma$ —lactone compound of the general formula I according to claim 1.according to claim 1 for the production of a pharmaceutical preparation—for the treatment of arteriosclerosis.

- 18. (Currently Amended) The A method according to claim 1 Use of using of wound healing comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to claim 1 according to claim 1 for the production of a pharmaceutical preparation for wound healing.
- 19. (Currently Amended) The A method according to claim 1 Use of using of treating a neoplastic disease comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted γ-lactone compound of the general formula I according to calim 1.according to claim 1 for the production of a pharmaceutical preparation for the treatment of neoplastic diseases.
- 20. (Currently Amended) The A method according to claim 1 Use of using of treating a fungal disease comprising administering to a patient a pharmaceutically effective amount of a pharmaceutical composition comprising at least one substituted \( \gamma \)-lactone compound of the general formula I according to claim 1. according to claim 1 for the production of a pharmaceutical preparation for the treatment of fungal diseases.